

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : A61K 31/585	A1	(11) International Publication Number: WO 94/08590 (43) International Publication Date: 28 April 1994 (28.04.94)
(21) International Application Number: PCT/US93/10063 (22) International Filing Date: 20 October 1993 (20.10.93) (30) Priority data: 07/963,469 20 October 1992 (20.10.92) US (60) Parent Application or Grant (63) Related by Continuation US 07/963,469 (CIP) Filed on 20 October 1992 (20.10.92) (71) Applicant (for all designated States except US): BTG PHARMACEUTICALS CORP. [US/US]; 1250 Broad- way, New York, NY 10001 (US).		(72) Inventor; and (75) Inventor/Applicant (for US only) : BERGER, Joseph, R. [US/US]; 6460 SW 109th Street, Miami, FL 33156 (US). (74) Agent: WHITE, John, P.; Cooper & Dunham, 30 Rocke- feller Plaza, New York, NY 10112 (US). (81) Designated States: AU, BR, CA, FI, HU, JP, KR, NO, NZ, PL, RU, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the</i> <i>claims and to be republished in the event of the receipt of</i> <i>amendments.</i>
(54) Title: A METHOD FOR AMELIORATING MUSCLE WEAKNESS/WASTING IN A PATIENT INFECTED WITH HUMAN IMMUNODEFICIENCY VIRUS-TYPE 1 (57) Abstract A method for attenuating the HIV-associated myopathy and muscle wasting associated with infection by human immu- nodeficiency virus-Type 1. Administration of oxandrolone in a daily dosage of about 2.5 to about 20 milligrams is described.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NE	Niger
BE	Belgium	GN	Guinea	NL	Netherlands
BF	Burkina Faso	GR	Greece	NO	Norway
BG	Bulgaria	HU	Hungary	NZ	New Zealand
BJ	Benin	IE	Ireland	PL	Poland
BR	Brazil	IT	Italy	PT	Portugal
BY	Belarus	JP	Japan	RO	Romania
CA	Canada	KP	Democratic People's Republic of Korea	RU	Russian Federation
CF	Central African Republic	KR	Republic of Korea	SD	Sudan
CG	Congo	KZ	Kazakhstan	SE	Sweden
CH	Switzerland	LJ	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovak Republic
CM	Cameroon	LU	Luxembourg	SN	Senegal
CN	China	LV	Latvia	TD	Chad
CS	Czechoslovakia	MC	Monaco	TG	Togo
CZ	Czech Republic	MG	Madagascar	UA	Ukraine
DE	Germany	ML	Mali	US	United States of America
DK	Denmark	MN	Mongolia	UZ	Uzbekistan
ES	Spain			VN	Viet Nam
FI	Finland				

A METHOD FOR AMELIORATING MUSCLE WEAKNESS/WASTING
IN A PATIENT INFECTED WITH
HUMAN IMMUNODEFICIENCY VIRUS-TYPE 1

5

Technical Field

The invention relates to the use of oxandrolone to attenuate myopathy and muscle weakness/wasting associated with infection by human immunodeficiency virus-Type 1.

10

Background of the Invention

Human immunodeficiency virus (HIV) associated myopathy and/or muscle weakness/wasting is a relatively common clinical manifestation of acquired immunodeficiency syndrome (AIDS). This is one of a number of neuromuscular disorders associated with the disease. There is some evidence to indicate that direct HIV infection of muscle may be at least partly responsible, occasionally resulting in a polymyositis-like disorder. In addition, zidovudine (AZT), an antiviral agent that is used widely in the clinical management of AIDS, has been associated with a toxic myopathy, presumably related to an inhibition of mitochondrial metabolism. In any event, the loss of muscle mass commonly observed in AIDS victims negatively impacts muscle function, however caused.

15

20

25

Individuals with HIV-associated myopathy or muscle weakness or wasting typically experience significant weight loss, generalized or proximal muscle weakness, tenderness, and muscle atrophy. Laboratory tests of samples from such individuals often reveal elevated levels of enzymes associated with muscle degeneration and necrosis, such as creatine kinase, aldolase, and aspartate amino transferase. Electromyographic test results for individuals with HIV-associated myopathy are typically consistent with

30

35

- 2 -

myopathic changes. Histopathologic tests may reveal muscle fiber necrosis associated with lymphocytic inflammatory infiltrates. In AZT myotoxicity, ragged red fibers are often observed.

5 Clinical management of HIV-associated myopathy and muscle weakness/muscle wasting varies. In individuals with AZT myopathy, withdrawal of this anti-retroviral agent may be associated with temporary improvement in strength and muscle bulk. Corticosteroid
10 therapy, such as the administration of prednisone, has been occasionally successful when inflammatory infiltrates have been detected in muscle. However, a potential drawback to this approach is that corticosteroids, because of their immunosuppressant
15 activity, may be harmful to individuals with AIDS who are already dangerously immunosuppressed as a consequence of the HIV infection.

 Furthermore, corticosteroid use itself is associated with myopathies and an increased
20 susceptibility to infections. Plasmapheresis has also been used with some success, although at least one patient has experienced, despite an increase in muscle strength, substantial weakness over a period of several weeks.

25

Summary of the Invention

 The present invention provides a method which employs oxandrolone (an anabolic steroid with weak androgenic activity) as an alternative approach to the
30 clinical management of HIV-associated myopathy/muscle weakness/muscle wasting. Loss in muscle mass (wasting) is attenuated, and body weight can be more readily maintained in this manner. Such an approach has been applied successfully to improve strength, reverse weight
35 loss, and provide an improved sense of well-being.

Importantly, no evidence of liver injury or other untoward side effects have been observed.

Oxandrolone preferably is administered orally; however, other routes of administration can be utilized as well.

The present method of ameliorating muscle weakness or muscle wasting in a patient infected with HIV-1 comprises administering to the patient daily a sufficient amount of oxandrolone to attenuate the patient's rate of muscle mass loss. To this end, oxandrolone may be administered, orally or otherwise, in a daily dose in the range of about 2.5 to about 20 milligrams. However, the response of individual patients may vary and in some instances a daily dose greater than 20 mg may be required to achieve the desired response. The daily dose may be divided into unit doses of about 1 to about 5 milligrams each, administered to the patient three times per day at about eight-hour intervals.

Detailed Description of the Preferred Embodiment

Oxandrolone (17-hydroxy-17-methyl-2-oxaandrostan-3-one) is a known compound that is commercially available. The preparation of oxandrolone is described, inter alia, in U. S. Patent No. 3,128,283 to Pappo, which description is incorporated herein by reference.

Pharmacologically, oxandrolone is a synthetic anabolic steroid similar in structure to testosterone, but having a different, lesser androgenic/anabolic activity ratio. In addition, oxandrolone is unique among all other testosterone analogues in that it contains an oxygen atom instead of a methylene group at the 2-position of the phenanthrene nucleus. In addition, oxandrolone lacks a 4-ene function in its A-ring. The anabolic potency of oxandrolone, estimated as

approximately 3 to 13 times that of testosterone, is believed to result from this unique structure.

5 Oxandrolone disposition and metabolism in man has been studied following oral administration of a 10 milligram dose. The study indicated that oxandrolone was rapidly and completely absorbed, yielding a mean peak plasma concentration of 417 micrograms of oxandrolone per milliliter at 66 minutes. The plasma concentration of oxandrolone declined in a biphasic
10 manner with a distribution half-life of approximately 30 minutes and an elimination half-life of 9.4 hours. Protein binding of oxandrolone was observed to be extensive.

15 In distinct contrast to other anabolic androgenic steroids such as methyltestosterone, fluoxymesterone, and micronized testosterone, oxandrolone taken orally is excreted mainly unchanged and unconjugated in urine. Urinary excretion of approximately 35 percent of an oral oxandrolone dose has
20 been observed within 72 hours after ingestion. After 96 hours, approximately 65 percent of the administered oxandrolone dose was excreted in urine. Fecal excretion accounts for less than about 3 percent over the same time period.

25 Oxandrolone compositions, upon administration in accordance with this invention, ameliorate myopathy and muscle weakness in patients suffering from infections by human immunodeficiency virus-Type 1. Anabolic steroids, as a class, are known to stimulate
30 appetite. Improved nutrition is important to individuals with AIDS who have experienced loss of lean body mass. Further, as a consequence of direct interaction with androgen and/or glucocorticoid receptors in muscle, anabolic steroids promote muscle

anabolism through both anabolic pathways and anticatabolic pathways.

5 Anabolic steroids, such as oxandrolone, also increase protein synthesis. For example, oxandrolone increased muscle protein synthesis in a study of acute uremic rats. Similarly, administration of oxandrolone preceded clinical improvement in appetite, cell mass, linear growth, and weight for height in boys with chronic renal failure. These observations are
10 consistent with anabolic activity. Oxandrolone may also stimulate the secretion of growth hormone and insulin-like growth factors.

 In addition to producing beneficial direct anabolic action, oxandrolone is also believed to act as
15 a delayed immunostimulant. In contrast, other appetite stimulants, such as dronabinol, that are currently under evaluation as appetite stimulants for AIDS patients can act as immunosuppressants in animals.

 For purposes of administration in accordance
20 with this invention, the active ingredient oxandrolone is combined with solid or liquid pharmaceutical carriers and formulated in unit dosage form using pharmacologically acceptable excipients, or dissolved or suspended in physiologically acceptable solvents or
25 liquid vehicles for oral, percutaneous, or topical administration.

 The overall daily dose of oxandrolone to provide a therapeutically effective amount in accordance with the method of this invention can be as low as about
30 2.5 milligrams and as high as about 20 milligrams, depending upon the patient's response and the mode of administration.

 The amount of the active ingredient within the aforementioned ranges that is to be administered depends
35 upon the age, weight and condition of the patient, as

well as on factors such as the frequency and route of administration. In formulating xandrolone, it is recognized that there may be differences between the immediate and the long term response. To account for these changes, the specific dosage given to a particular patient is based also on the individual patient's response. Preferably, oxandrolone is orally administered to the patient daily for a time period in the range of about 2 weeks to about 6 months.

Attenuation of the rate of muscle mass loss in a patient can be ascertained by comparing the patient's rate of weight loss before oxandrolone therapy with that after the administration of oxandrolone has been commenced. Alternatively, or in addition, the patient's urinary nitrogen level can be monitored, a well-known expedient. A decrease in the patient's urinary nitrogen level is indicative of a decrease in muscle mass loss.

Similarly, the maintenance of a relatively stable patient's total body potassium level, as well as an increase in the patient's total body potassium level, upon oxandrolone administration indicates that a therapeutically effective amount of oxandrolone is being administered. A patient's total body potassium level can be monitored, for example, as described in Kotler et al., *The American Journal of Clinical Nutrition*, 42:1255-1265 (December 1985) and Pierson, Jr., et al., *Am. J. Physiol.*, 246 (Renal Fluid Electrolyte Physiol. 15):F234-F239 (1984).

The route of administration can be oral, percutaneous, transdermal, sublingual, buccal, intravenous, intramuscular, or the like. Of these, oral administration is preferred. The patient's daily dose of the active ingredient preferably is in the range of about 7.5 milligrams, but may exceed 20 milligrams based on clinical response. This daily dose can be given in

tablet form as a single dose, or as plural divided doses, preferably 2 to 3 divided doses. The requisite daily dose can also be supplied continuously, for example, by a transdermal patch worn by the patient or
5 intravenously. If the oxandrolone is administered orally, dosages in the range of about 2 to about 5 milligrams three to four times daily typically may be utilized.

Oxandrolone tablets are manufactured using
10 standard solid dose form technology in accordance with United States Pharmacopeia (USP) specifications (see, for example, The United States Pharmacopeia, 22nd Revision, pp. 981-982). Specifically, a typical 150-milligram tablet contains the following:

15	Oxandrolone, USP	2.5 mg
	Corn Starch, NF	30.0 mg
	Lactose NF (hydrous)	113.0 mg
	Hydroxypropyl Methylcellulose, USP	3.0 mg
	Magnesium Stearate	<u>1.5 mg</u>
20		150.0 mg

The terms "unit dosage form" and "unit dose" as used in the present specification and claims refer to a physically discrete unit or units suitable as unitary doses for patients, each unit containing a predetermined
25 quantity of the active ingredient calculated to produce the desired therapeutic effect in association with the pharmacologically acceptable carrier. The specifications for the unit dosage forms of this invention are dictated in part and are also dependent
30 upon (a) the unique characteristics of the active ingredient and (b) the particular therapeutic effect to be achieved, as well as upon limitations inherent in the art of compounding such active ingredient for the therapeutic use disclosed in detail in this
35 specification. Examples of suitable unit dosage forms

in accordance with this invention are tablets, pills, powder packets, wafers, cachets, segregated multiples of any of the foregoing, transdermal patches, aliquots of injectables, and the like forms.

5 The primary response variables are patient's total body potassium, body weight, muscle mass, muscle strength, improvement in or increased appetite, and general sense of well-being. In addition, improvement
10 in immune status (or at a minimum, no worsening of immune function) in response to oxandrolone is significant as well.

 An important question regarding the use of any drug in combination with anti-retroviral therapy is whether drug interactions may occur that would diminish
15 AZT efficacy or increase the frequency of severity of AZT-related adverse reactions. TABLE 1 compares various published pharmacological parameters for oxandrolone and AZT and illustrates important differences between the
20 two drugs.

TABLE 1
Comparison of Selected Oxandrolone
and AZT Pharmacology Parameters

5	<u>Parameter</u>	<u>Oxandrolone</u>	<u>AZT</u>
	Oral Bioavailability	100%	65%
	Tmax	1.1 hr	0.7 hr
10	Biological T1/2	9.4 hr	1.1 hr
	Vd	578 ml/kg	>1400 ml/kg
15	Protein Binding	>95%	25-35%
	Plasma Clearance	43 ml/kg/hr	>1300 ml/kg/hr
	Metabolism	Little	Extensive
20	Glucuronidation	Little	Substantial
	Urinary Excretion	Extensive; primarily parent compound	Extensive; parent and glucuronide conjugated
25	Target Organ Toxicity	Liver (anabolic steroids as a class)	Hematopoietic system (e.g., anemia, granulocytopenia)
30	Known Drug Interactions	Anticoagulants; oral hypoglycemic agents; adrenal steroid when edema present	Drugs that may: (a) inhibit glucuronidation (e.g., aspirin, acetaminophen) or urinary excretion (e.g., probenecid); (b) adversely affect blood cell number and function; and (c) nephrotoxic or cytotoxic
35			

- 10 -

Because oxandrolone is primarily protein bound, whereas AZT is primarily non-protein bound, oxandrolone will not compete appreciably with AZT for binding sites in plasma. Consequently, administration of oxandrolone to patients on AZT therapy is unlikely to alter the level of free AZT in the blood. Likewise, the administration of AZT is unlikely to alter the level of free oxandrolone in the blood. An oxandrolone-AZT drug interaction involving binding site displacement is, therefore, extremely unlikely.

AZT is rapidly metabolized and excreted in the urine--a significant quantity is excreted in the form of glucuronide conjugates. In sharp contrast, oxandrolone, perhaps due to presence of a lactone group and the absence of a 4-ene function in the A-ring, undergoes little hepatic metabolism and is excreted primarily unchanged and unconjugated in urine. Thus, in contradistinction to other drugs that may competitively inhibit glucuronidation and thereby potentially slow the rate of AZT metabolism, such as aspirin, acetaminophen, or indomethacin, the present active agent, oxandrolone, is not believed to affect AZT metabolism.

Furthermore, oxandrolone is neither nephrotoxic nor cytotoxic. Accordingly, oxandrolone is not expected to interfere with the renal excretion of AZT or its metabolites. To the contrary, oxandrolone has been safely and effectively used in patients with chronic renal disease to stimulate growth and increase lean body mass. In well-controlled studies of oxandrolone for the clinical management of critically ill patients with acute alcoholic hepatitis, oxandrolone administered at daily doses of up to 80 mg/day for four weeks and 40 mg/day for eight weeks did not result in any drug-related nephrotoxicity.

- 11 -

While it is known that anabolic androgenic steroids have been associated with potentially life-threatening forms of liver disease, including peliosis hepatitis, cholestatic jaundice, and hepatocellular neoplasms, specific reports in the medical literature regarding liver disease in oxandrolone-treated patients, at the dosages proposed for use in the clinical management of HIV associated muscle weakness/wasting (i.e., about 2.5 to about 20 mg/day) are rare.

Oxandrolone and AZT have different mechanisms of action. They also function in different sites of cellular action at the receptor level. Oxandrolone functions via interaction with androgen and glucocorticoid receptors, whereas AZT, once phosphorylated, acts to inhibit HIV reverse transcription. Thus, competitive inhibition of AZT by oxandrolone at the cellular level also is considered unlikely.

Neither has oxandrolone been associated with anemia or granulocytopenia, two frequently occurring and potentially serious side effects associated with AZT therapy. To the contrary, anabolic androgenic steroids have been used clinically to stimulate erythropoiesis in hypoanemias, aplastic anemias, hemolytic anemias, renal anemias, anemias due to cytotoxic therapy, and various leukemias. It has been reported recently that androgens augment beneficial effects of erythropoietin in the treatment of anemia resulting from end-stage renal disease.

Data derived from animal models and human clinical studies indicate that anabolic steroids are unlikely to suppress immune function in patients infected with HIV. For example, anabolic steroids can stimulate granulopoiesis in mice, as evidenced by stimulation of granulocytic colony-forming cells derived

from spleen and bone marrow. Similarly, an anabolic steroid known as nandrolone decanoate enhanced macrophage activity and cell-mediated immunity in patients with uterine cervical cancer when administered parentally. In related studies, anabolic steroids increased peripheral lymphocyte and monocyte counts, Immunoglobulin G (IgG) levels, and PHA-blastoid transformation of peripheral lymphocytes. In those studies, β_2 -microglobulin levels simultaneously decreased.

IgG is one of a class of antibodies secreted by B cells (i.e., B-lymphocytes) in response to an antigenic challenge (e.g., foreign protein like that from bacteria). In the case of HIV infection, humoral immune function (i.e., B-cell mediated) is significantly impaired. Accordingly, when HIV-infected individuals are challenged with a specific antigen, the typical response of B-cell proliferation, differentiation and secretion of antibodies (e.g., IgG) is diminished or absent. This decline in humoral immune function coupled with defects in cellular immune (i.e., T-cell) function contributes to the overall failure of the immune system to respond in an appropriate manner to challenge. B-cells in AIDS victims are, by mechanisms unknown, hyperstimulated to secrete large amounts of immunoglobulins that make the humoral system refractory to new antigens. The result is that the patient's system no longer recognizes new antigens and does not respond.

In animal studies in which anabolic steroids have been reported to increase IgG and PHA-blastoid activity, these changes occurred as a result of immune system stimulation, and are positive responses. β -microglobulin is a cell surface protein that is found on all nucleated cells and it is released into the serum

- 13 -

during cell turnover. Generally, β -microglobulin is considered a marker of infectious, inflammatory, malignant and autoimmune disease activity. In several AIDS studies, β -microglobulin levels correlated with disease progression and T4 (T-helper) cell counts. In the case of therapy with oxandrolone, for example, a decrease in β -microglobulin levels is desirable. Thus, animal data showing reduced plasma levels of β -microglobulin in response to anabolic steroids is evidence of a positive effect and suggestive of similar activity in man.

Accordingly, there are no reasons to believe that the administration of an anabolic steroid in general and oxandrolone in particular would have adverse effects on the immune system. Generally, the target organ of toxicity for these drugs is the liver--probably because this is where most are metabolized. Oxandrolone, however, has a remarkably good safety profile in man as a likely consequence of its resistance to hepatic metabolism; an oral dose is excreted primarily in urine as the parent compound, as stated hereinabove.

Data from clinical trials in patients with severe alcoholic liver disease provide further evidence that oxandrolone is not likely to suppress immune function in patients with HIV infection. Ethanol abuse is associated with loss of lymphocyte functions, particularly T-cell dependent immune responses. Previous researchers have observed that oxandrolone significantly improved lymphocyte number in patients with severe alcoholic hepatitis. Because the loss of lymphocytic function by alcoholic liver disease parallels, to a significant degree, the loss of T-cell function due to HIV infection, it is reasonable to

- 14 -

hypothesize that oxandrolone will increase the T-Cell function of HIV-infected patients.

Therefore, these data from laboratory animals and human studies indicate that suppression of the immune system by anabolic steroids, such as oxandrolone, is unlikely. Nonetheless, subjects undergoing oxandrolone therapy, as a precaution, should be monitored for changes in lymphocyte number, particularly CD4+ and CD8+, as is routinely done for patients who undergo steroid therapy.

In summary, based on the differences between AZT and oxandrolone with respect to pharmacokinetics, metabolism, reported drug interactions, mechanisms of action, and reported toxicities, oxandrolone and AZT can be safely used in combination for subjects infected with the Type-1 HIV virus and suffering from HIV-associated myopathy. The use of oxandrolone in patients on AZT therapy is, on the basis of known drug interactions, also consistent with current FDA-approved labeling for AZT and oxandrolone.

The following example demonstrates the effectiveness of oxandrolone in attenuating the effects of HIV-associated muscle weakness or muscle wasting in an AIDS patient.

25

EXAMPLE

A patient, a thirty-two year old homosexual man, known to be HIV-seropositive since February 1989, noted difficulty opening drawers and bottles in May 1989. The patient weakened progressively and, during a physical examination in September 1989, demonstrated by confrontation testing the weakness of neck flexion and proximal limbs. However, his muscle stretch reflexes remained normal. Laboratory tests showed the patient's creatine kinase level to be 286 International Units per

- 15 -

liter, much higher than the normally observed range for creatine kinase of about 40-200 Units per liter.

5 Zidovudine (azidothymidine or AZT) was initiated at 500 milligrams daily, but the patient's strength continued to decline through February 1990. He complained of an inability to ascend a flight of stairs. The patient exhibited greater weakness and atrophy of neck flexors and extremity muscles during another physical examination performed at this time. An
10 electromyogram revealed a decrease of amplitude and duration of the patient's motor unit potentials and increased recruitment in selected muscles of his right upper extremity. The patient's creatine kinase tested at 456 Units per liter. A muscle biopsy revealed
15 numerous myofibers, abundant ragged red fibers, and numerous eosinophilic inclusions. Round cell inflammatory infiltrates were also noted. In light of these developments, the zidovudine treatment was terminated.

20 Substantial improvement initially followed the discontinuation of zidovudine. However, because of a subsequent continued and progressive weakness rendering it difficult for the patient to ascend or descend a flight of stairs, a prednisone therapy (60 mg daily) was
25 initiated. No significant improvement accompanied the use of prednisone.

 Thereafter, a trial period of oral oxandrolone administration (2.5 milligrams, three times daily, in tablet form) was initiated. Within two weeks of the
30 initiation of the oxandrolone therapy, the patient noted an improved sense of well being, became stronger, and gained weight. Within one month, he was able to ascend and descend stairs without problems. Confrontation testing revealed nearly normal strength. The patient's
35 weight increased from 115 pounds to 130 pounds. The

- 16 -

patient's muscle atrophy was alleviated as well. Liver functions were closely monitored for signs of elevation, but undesirable side effects were not detected.

5 After several months of the aforementioned therapy with oxandrolone, the patient was no longer able to obtain oxandrolone for use as a medication. Weakness and weight loss ensued. Trials of other anabolic preparations, specifically stanazol and oxymethalone, did not return the patient to his previous levels of
10 function and strength.

The EXAMPLE demonstrates that oxandrolone can be a beneficial alternative for clinical management of HIV-associated myopathy and muscle weakness and wasting.

15 It is intended that the foregoing description is by way of illustration only and is not to be construed as limiting the invention in any way except in the spirit and scope of the appended claims.

WHAT IS CLAIMED IS:

1. Use of oxandrolone in the manufacture of a composition for the amelioration of myopathy and muscle weakness in an amount effective to attenuate the rate of muscle mass loss in a patient infected with a Type-1 human immunodeficiency virus.
2. The use of oxandrolone according to claim 1 wherein the oxandrolone is administered to the patient in a daily dosage in the range of about 2.5-30 milligrams.
3. The use of oxandrolone according to claim 2 wherein the oxandrolone is administered to the patient in a daily dosage of about 7.5 milligrams.
4. The use of oxandrolone according to claim 2 wherein the oxandrolone is administered to the patient in a daily dosage of about 15 milligrams.
5. The use of oxandrolone according to claim 1 wherein the oxandrolone is administered to the patient as a unit dose of about 1-5 milligrams 3 times a day at about eight hour intervals.
6. The use of oxandrolone according to claim 1 wherein the resulting composition is administered percutaneously.
7. The use of oxandrolone according to claim 1 wherein the resulting composition is administered intravenously.
8. The use of oxandrolone according to claim 1 wherein the resulting composition is administered intramuscularly.
9. The use of oxandrolone according to claim 1 wherein the resulting composition is administered sublingually.

-18-

10. The use of oxandrolone according to claim 1 wherein the resulting composition is administered transdermally.
- 5 11. The use of oxandrolone according to claim 1 wherein the resulting composition is administered orally.
12. The use of oxandrolone according to claim 11 wherein the resulting composition is in the form of a tablet.
- 10 13. The use of oxandrolone according to claim 1 wherein the resulting composition may be administered for a time period in the range of about 2 weeks to about 6 months.
- 15 14. Use of oxandrolone in the manufacture of a composition for the amelioration of myopathy and muscle weakness in a patient infected with a Type-1 human immunodeficiency virus such that the resulting composition may be an oral composition suitable for administration for a time period in the range of about 2 weeks to about 6 months.
- 20 15. A pharmaceutical composition comprising oxandrolone in an amount effective to attenuate the rate of muscle mass loss for the amelioration of myopathy and muscle weakness in a patient infected with a Type-1 human immunodeficiency virus and a pharmaceutically acceptable carrier.
- 25 16. The composition of claim 15 wherein the effective amount provides a daily dosage in the range of about 2.5 to about 30 milligrams oxandrolone.
- 30 17. The composition of claim 15 wherein the effective amount provides a daily dosage of about 7.5 milligrams oxandrolone.
- 35

18. The composition of claim 15 wherein the effective amount provides a daily dosage of about 15 milligrams oxandrolone.
- 5 19. The composition of claim 15 wherein the effective amount provides a unit dose of about 1 to about 5 milligrams oxandrolone and is administered 3 times per day at about equally spaced intervals.
- 10 20. The composition of claim 15 for percutaneous administration.
21. The composition of claim 15 for intravenous administration.
- 15 22. The composition of claim 15 for intramuscular administration.
23. The composition of claim 15 for sublingual administration.
- 20 24. The composition of claim 15 for transdermal administration.
- 25 25. The composition of claim 15 for oral administration.
26. The composition of claim 25 in the form of a tablet.
27. The composition of claim 15 for administration over a time period in the range of about 2 weeks to about 6 months.
- 30 28. A composition comprising oxandrolone for the amelioration of myopathy and muscle weakness in a patient infected with a Type-1 human immunodeficiency virus and a pharmaceutically acceptable carrier such
- 35

that the composition is an oral composition and is appropriate for administration for a time period in the range of about 2 weeks to about 6 months.

- 5 29. A method for ameliorating myopathy and muscle weakness
in a patient infected with a Type-1 human
immunodeficiency virus which comprises administering to
said patient oxandrolone in an amount sufficient to
attenuate the rate of muscle mass loss in said patient.
- 10 30. The method in accordance with claim 29 wherein the
oxandrolone is administered to said patient in a daily
dosage in the range of about 2.5 to about 30
milligrams.
- 15 31. The method in accordance with claim 29 wherein the
daily dosage of the oxandrolone is about 7.5
milligrams.
- 20 32. The method in accordance with claim 29 wherein the
daily dosage of the oxandrolone is about 15 milligrams.
- 25 33. The method in accordance with claim 29 wherein the
oxandrolone is administered to said patient as a unit
dose of about 1 to about 5 milligrams three times per
day at about eight-hour intervals.
- 30 34. The method in accordance with claim 29 wherein the
oxandrolone is administered percutaneously.
- 35 35. The method in accordance with claim 29 wherein the
oxandrolone is administered intravenously.
36. The method in accordance with claim 29 wherein the
oxandrolone is administered intramuscularly.

37. The method in accordance with claim 29 wherein the oxandrolone is administered sublingually.
38. The method in accordance with claim 29 wherein the oxandrolone is administered transdermally.
39. The method in accordance with claim 29 wherein the oxandrolone is administered orally.
40. The method in accordance with claim 39 wherein the oxandrolone is administered in the form of a tablet.
41. The method in accordance with claim 29 wherein administration is continued over a period in the range of about 2 weeks to about 6 months.
42. A method for ameliorating HIV-associated myopathy and muscle wasting in a patient infected with a Type-1 human immunodeficiency virus which comprises orally administering a therapeutically effective amount of oxandrolone to said patient daily for a time period in the range of about 2 weeks to about 6 months.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US93/10063

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) :A61K 31/585

US CL :514/175

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/175

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
none

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS, APS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Chemical Abstracts, Volume 74, no. 4, issued 17 May 1991, (Columbus Ohio, USA)), Bourse et al, "Comparison of the androgenic and myotrophic activities of some anabolic steroids in the castrated rat.", see abstract no. 72305d, (Res. Div., Hoffman-La Roche Inc., Nutley, N.J.) J. Steroid Biochem., 1970, vol.1 (4), pp. 349-54.	1-42



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	* T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
* A* document defining the general state of the art which is not considered to be part of particular relevance	* X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
* E* earlier document published on or after the international filing date	* Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
* L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	* A*	document member of the same patent family
* O* document referring to an oral disclosure, use, exhibition or other means		
* P* document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

03 FEBRUARY 1994

Date of mailing of the international search report

17 FEB 1994

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Authorized officer

RUSSELL TRAVERS

Facsimile No. NOT APPLICABLE

Telephone No. (703) 308-1235

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US93/10063

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Chemical Abstracts, Volume 115, no. 15, Issued 02 July 1991 (Columbus Ohio, U.S.A.), Aroonsakul, Chaivane, "Method of treatment of central nervous system diseases such as Alzheimer's disease and Parkinson's disease and method of diagnosing Alzheimer's disease", see abstract no. 151902q, Pat. Specif. (Aust.) AU 604,820 (Cl. A61K 31/565), 03 Jan. 1991, Appl. 88/10,125, 08 Jan 1988, 34pp.	1-42
Y	Chemical Abstracts, Volume 107, no. 18, Issued 27 September 1987, (Columbus Ohio, U.S.A.), Lone et al, "The use of oxandrolone, an anabolic-androgenic steroid, as a growth promotant in carp. Effect on tissues molecular growth responses", see abstract nol 174950c, (Dep. Biol. Sci., Univ. Aston, Birmingham, UK B4 7ET). Pak J. Zool. 1986, vol. 18(2), pp. 153-69.	1-42